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I CLAIM: A method for monitoring biological micro-structure activity which 1. produces detectable signals characterizing events, comprising the steps of estimating the

fundamental frequency of the occurrence of events from the detectable signals, without

detecting the occurrence of individual events.

The method of claim 1 wherein events are analyzed during an analysis 2. window which spans more than one event.

The method of any of claims 1 or 2 wherein the events are signals 3. produced by biological micro-structures which are displaced from their original environment.

The method of any of claims 1 or 2 wherein the events are signals 4. produced by living cells.

The method of claim 4 wherein the events are signals produced by the 5. Islets of Langerhans.

The method of any one of claims 10-2 wherein the estimating step 6. includes an autocorrelation operation.

The method of claim 6 further including one of the following steps: 7. estimating the fundamental frequency based upon a lower autocorrelation value disposed among several adjacent peaks;

treating "unvoiced" segments of the detectable signal as undecided as to pitch and estimating the pitch of those segments through subsequent processing;

seeking to estimate the fundamental frequency in the range of .25 to 5

Hertz;

utilizing an analysis window duration in the range of several seconds; performing a pre-processing operation which has the effect of increasing the effective duration of an event; and

utilizing an autocorrelation process which performs segmented

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12		autocorrelation.
1		8. The method of Claim 1 further comprising the step of using the number
2		of "unvoiced" windows within a predetermined time as a measure of a blood constituent level
3		of a patient.
1		9. The method of Claim 1 further comprising the step of using a sureness
2		grade as measure of a blood or tissue constituent level of a patient.
1		10. The method of Claim 1 further comprising the step of using the
2		fundamental frequency as a measure of a blood or tissue constituent level of a patient.
1		11. The method of Claim 1 wherein the blood constituent level is the blood
2 1 2 2 1 2 2 1 2 1 2 1 2 1 1 2 1 1 1 1		glucose level in the vicinity of the biological micro-structure.
1		12. The method of claim 1 wherein the events are electrical signals
2 3		produced by living cells in the Islets of Langerhans used as a probe within a patient and the
		fundamental frequency estimate is used as a measure of blood glucose level of the patient.
3 1	Ø	13. The method of any one of claims 2, wherein an analysis window spans
2		a duration of up to 40 times the interval between successive events
1		14. In a system for monitoring biological micro-structure activity which
2		produces detectable signals characterizing events, a sensor capable of receiving the sensible
3		signals and a processor including a module for estimating the fundamental frequency of the
4		occurrence of events from the detectable signals, without first detecting the occurrence of
5		individual events.
6		15. The system of claim 14 wherein the processor further comprises a
7		module for producing an analysis window during which events are analyzed, the analysis
8		window spanning more than one event.

The system of Claim 14 or 15 wherein the module for estimating

1			includes components	to perform an autocorrelation operation.
1			17.	The system of Claim 16 wherein the module for estimating further
2	2		includes one of the f	ollowing submodules:
. 3	3			a submodule which estimates the fundamental frequency based upon
4	Ļ		a lower autocorrelati	on value disposed among several adjacent peaks;
5	5		1	a sub-module which identifies "unvoiced" segments of the detectable
ϵ	3		signal as undecided	as to pitch, the pitch of those segments being estimated by a subsequent
7	7		processing submodu	le;
8	3			a submodule which controls the estimate of the fundamental frequency
)		to be in the range of	.25 to 5 Hertz;
40)		·	a submodule controlling the analysis window to have a duration in the
	l		range of several second	onds; and
72	2			a submodule performing a pre-processing operation which has the effect
4	3		of increasing the effe	ective duration of an event.
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13 <i>.</i>		0	18.	The system of Claim 14 or 15 wherein the processor is constructed to
	2		perform a segmented	autocorrelation process.
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IJ.	1	F	19.	The system of claim 14 or 15 wherein the sensor is a probe capable of
:	2			itted by living cells in the Islets of Langerhans, the frequency estimate
;	3		being an indication	f blood glucose level of a patient in which those cells are present.
				claim 15
	1	α	20.	The system of any one of elaims 15, wherein an analysis window spans
	2		a duration of up to 4	times the interval between successive events.
		12	ai>	
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